=> d his

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(FILE 'HOME' ENTERED AT 03:10:17 ON 28 JAN 2002)
     FILE 'REGISTRY' ENTERED AT 03:10:29 ON 28 JAN 2002
L1
               STRUCTURE UPLOADED
L2
             46 S L1
L3
           1009 S L1 FULL
     FILE 'CA' ENTERED AT 03:11:49 ON 28 JAN 2002
L4
           454 S L3
L5
            445 S L4 AND PD < NOVEMBER 2000
             7 S L5 AND ANTIMYCOBACTERIAL?
L6
     FILE 'CAOLD' ENTERED AT 03:14:20 ON 28 JAN 2002
L7
           270 S L3
     FILE 'REGISTRY' ENTERED AT 03:14:50 ON 28 JAN 2002
              E 13410-86-1/RN
L8
              1 S E3
               E 2779-55-7/RN
L9
              1 S E3
              E 149-17-7/RN
L10
             1 S E3
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LOGINID:ssspta1612BXR
PASSWORD:
TERMINAL (ENTER 1, 2, 3, OR ?):2

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Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

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=>
Uploading 9699732a.str

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

Structure attributes must be viewed using STN Express guery preparation.

=> s 11

SAMPLE SEARCH INITIATED 03:11:35 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED - 374 TO ITERATE

100.0% PROCESSED 374 ITERATIONS

46 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 6320 TO 8640 514 TO PROJECTED ANSWERS: 1326

46 SEA SSS SAM L1

=> s l1 full

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FULL SEARCH INITIATED 03:11:44 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 7960 TO ITERATE

100.0% PROCESSED 7960 ITERATIONS 1009 ANSWERS SEARCH TIME: 00.00.01

1009 SEA SSS FUL L1

=> file ca

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SINCE FILE TOTAL ENTRY SESSION 140.54 140.69

FULL ESTIMATED COST

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FILE COVERS 1907 - 24 Jan 2002 VOL 136 ISS 5 FILE LAST UPDATED: 24 Jan 2002 (20020124/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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The CA file now provides online access to patents and literature covered in CA from 1907 to the present. Bibliographic information and abstracts were added in 2001 for over 3.8 million records from 1907-1966.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

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=> s 13

L5

L4 454 L3

=> s 14 and pd < november 2000

20047988 PD < NOVEMBER 2000 (PD<20001100) 445 L4 AND PD < NOVEMBER 2000

=> s 15 and antimycobacterial?

880 ANTIMYCOBACTERIAL? 7 L5 AND ANTIMYCOBACTERIAL?

=> d 16, ibib abs fhitstr, 1-7

L6 ANSWER 1 OF 7 CA COPYRIGHT 2002 ACS ACCESSION NUMBER: 132:329080 CA

TITLE: Isoniazid-related copper(II) and nickel(II) complexes

with antimycobacterial in vitro activity.

Part 9

AUTHOR(S): Bottari, B.; Maccari, R.; Monforte, F.; Ottana, R.;

Rotondo, E.; Vigorita, M. G.

CORPORATE SOURCE: Dipartimento Farmaco-chimico, Facolita di Farmacia,

Universita di Messina, Messina, 98168, Italy

SOURCE: Bioorg. Med. Chem. Lett. (2000), 10(7),

657-660

CODEN: BMCLE8; ISSN: 0960-894X

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Isonicotinoylhydrazones (HL) obtained from the primary antituberculosus agent Isoniazid were used as monoanionic ligands (L) to prep. Cu(II) and Ni(II) octahedral complexes [ML2(H2O)2]. Their antimycobacterial in vitro activity was evaluated against M. tuberculosis H37Rv in comparison with the ligands. Some complexes displayed MIC values .ltoreq.

0.2 .mu.g/mL.

IT **86189-87-9**

RL: RCT (Reactant)

(for prepn. of copper or nickel isonicotinoylhydrazone complexes)

RN 86189-87-9 CA

CN 4-Pyridinecarboxylic acid, [(4-fluorophenyl)methylene]hydrazide (9CI)

(CA

L6

INDEX NAME)

REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR

THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L6 ANSWER 2 OF 7 CA COPYRIGHT 2002 ACS ACCESSION NUMBER: 132:205321 CA

TITLE: Minimal inhibitory concentration and minimal

bactericidal concentration determination of

isonicotinic acid derivatives against Mycobacterium

tuberculosis

AUTHOR(S): Sato, D. N.; Bacha, C. T. M.; Garibott, D.; Bottcher,

M.; Errera, M. C.; Presotto, P.; Melles, Carmo Elias

Page 5

Andrade

CORPORATE SOURCE:

Instituto Adolfo Lutz, Laboratorio I de Ribeirao

Preto, Brazil

SOURCE:

Rev. Inst. Adolfo Lutz (1999), 58(1), 25-29

CODEN: RIALA6; ISSN: 0073-9855

PUBLISHER:

Instituto Adolfo Lutz

DOCUMENT TYPE:

Journal

LANGUAGE:

Portuguese

AB Tuberculosis still remains as a worldwide public health problem with high morbidity and mortality in developing countries. The increase of strains of M. tuberculosis that are resistant to antimycobacterial agents is a worldwide problem. Consequently, it is urgently necessary to develop antimycobacterial drugs which are more effective that those used in conventional treatment of tuberculosis. Twelve

isonicotinic

acid derivs. were evaluated for in vitro activity against M. tuberculosis H37Ra-ATCC 25177. The MIC and minimal bactericidal concn. (MBC) of M. tuberculosis H37Ra was detd. by broth macrodilution method. -He MIC of all derivs. showed a range of 0.062~0.250~mu.g/mL. In general, the MBC values for all derivs. were 2-fold higher than their corresponding MICs values. These MICs and MBCs values are close to isoniazid, considered

the

gold std. in this study.

ΙT 93-47-0

> RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (inhibitory activity of isonicotinic acid derivs. against

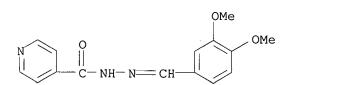
Mycobacterium

tuberculosis)

RN 93-47-0 CA

CN 4-Pyridinecarboxylic acid, [(3,4-dimethoxyphenyl)methylene]hydrazide (9CI)

(CA INDEX NAME)



ANSWER 3 OF 7 CA COPYRIGHT 2002 ACS

ACCESSION NUMBER:

132:12262 CA

TITLE:

Polyether hydrazines and hydrazones as selective

antimycobacterial agents

INVENTOR(S):

Chupakhin, Oleg Nikolaevich; Fedorova, Olga

Vasilievna; Rusinov, Gennady Leonidovich; Mordovskoi, Georgy Georgievich; Khomenko, Alexandr Grigorievich; Golyshevskaya, Valentina Ivanovna; Zueva, Marina

Nikolaevna; Ovchinnikova, Irina Georgievna

PATENT ASSIGNEE(S):

Institut Organicheskogo Sinteza Uralskogo Otdeleniya

Rossiiskoi Akademii Nau, Russia

SOURCE:

PCT Int. Appl., 30 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent Russian

LANGUAGE:

יית. 1 ייתו

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE WO 9959961 19991125 A1 WO 1999-RU165 19990518 <--W: CA, CN, JP, KR, US RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE RU 2137750 19990920 RU 1998-109494 C1 19980519 <--EP 1081131 EP 1999-922691 Α1 20010307 19990518 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE PRIORITY APPLN. INFO.: RU 1998-109494 A 19980519 WO 1999-RU165 W 19990518

OTHER SOURCE(S):

MARPAT 132:12262

GΙ

AB Title compds. such as I, and some of their metal complexes, were prepd. and tested as antimycobacterial agents. Thus, a suspension of 0.94 g 1,5-bis(2-formylphenoxy)-3-oxapentane and 0.83 g isonicotinic acid hydrazide in 20 mL 80% ethanol was subjected to ultrasound irradn. (22 kHz) for 3-5 min to give a 95% yield of I. The products showed high antimycobacterial activity against several strains in vitro; one compd. was also tested in guinea pigs.

IT 251364-96-2P
 RL: BAC (Biological activity or effector, except adverse); RCT
(Reactant);

Ι

SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(polyether hydrazines and hydrazones as selective antimycobacterial agents)

RN 251364-96-2 CA

CN 4-Pyridinecarboxylic acid, [oxybis(2,1-ethanediyloxy-2,1-phenylenemethylidyne)]dihydrazide (9CI) (CA INDEX NAME)

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L6 ANSWER 4 OF 7 CA COPYRIGHT 2002 ACS

ACCESSION NUMBER: 124:50481 CA

TITLE: 2-(4-Pyridyl)-.DELTA.2-1,3,4-oxadiazolines from

isonicotinoylhydrazones and diazomethane as potential

antimycobacterial and anti-HIV agents. V

AUTHOR(S): Vigorita, Maria Gabriella; Ottana, Rosaria; Zappala,

Carmela; Maccari, Rosanna; Pizzimenti, Francesco C.;

Gabbrielli, Gabriele

CORPORATE SOURCE: Facolta Farmacia, Univ. Messina, Messina, 98168,

Italy

SOURCE: Farmaco (1995), 50(11), 783-6

6

CODEN: FRMCE8

DOCUMENT TYPE: Journal

LANGUAGE: English

Me N-N

AB The 5-aryl-4-methyl-2-(4-pyridyl)-.DELTA.2-1,3,4-oxadiazolines, previously

synthesized alòng with isomer

4-aryl-1-methoxy-1-(4-pyridyl)-2,3-diaza-1,3-

Ι

butadienes from benzaldehyde isonicotinoylhydrazones and diazomethane, were tested for in vitro activity against both Mycobacterium tuberculosis and some atypical mycobacterial strains as well as against human immunodeficiency virus (HIV-1). Some halophenyl derivs.(I (Ar=3-chlorophenyl), II (Ar=3-triflouromethylphenyl), III (Ar=2,4-dichlorophenyl), and IV (Ar=3,4-dichlorophenyl)) were found to display MIC ranges from 1 to 10 .mu.g/mL against H 37 Rv and a clin. isolate tubercular strain, whereas against M. avium (MAC) the MICs were higher than 20 .mu.g/mL. When the combinations of oxadiazolines with

ethambutol, acting as an inhibitor of cell wall synthesis, were assayed

on

MAC strain a synergistic effect was demonstrated for trifluoromethyl derivs. The antimycobacterial profiles of 4-aryl-1-methoxy-1-(4pyridyl)-2,3-diaza-1,3-butadienes and 5-aryl-4-methyl-2-(4-pyridyl)-.DELTA.2-1,3,4-oxadiazolines analogs are compared and discussed. As is the case for 4-aryl-1-methoxy-1-(4-pyridyl)-2,3-diaza-1,3-butadienes, nosubstantial anti-HIV in vitro activity was found in selected .DELTA.2-oxadiazolines; a moderate cytotoxicity, however, appears to be a common property.

ΙT 144293-93-6

RL: RCT (Reactant)

(reaction with diazomethane)

144293-93-6 CA RN

CN 4-Pyridinecarboxylic acid, [(2-fluorophenyl)methylene]hydrazide (9CI)

(CA

INDEX NAME)

ANSWER 5 OF 7 CA COPYRIGHT 2002 ACS

ACCESSION NUMBER:

122:314426 CA

TITLE:

Halogenated isoniazid derivatives as possible

antimycobacterial and anti-HIV agents - III

AUTHOR(S): Vigorita, Maria Gabriella; Ottana, Rosaria; Zappala, Carmela; Maccari, Rosanna; Pizzimenti, Francesco C.;

Gabbrielli, Gabriele

CORPORATE SOURCE:

Dip. Farm.-Chim., Fac. Farm., Messina, 98168, Italy

SOURCE:

Farmaco (1994), 49(12), 775-81

CODEN: FRMCE8

DOCUMENT TYPE:

LANGUAGE:

Journal English

GΙ

$$_{R1}$$
 $_{CH=NN=C(OMe)}$
 $_{N}$

Acetophenone isonicotinoylhydrazones (I, R = H, 2-Cl, 3-F, etc.; R1 =AΒ 2-F,

4-OMe, etc.) and 4-aryl-1-methoxy-1-(4-pyridyl)-2,3-diaza-1,3-butadienes (II, same R = H, 3-Cl, etc.; R1 = H, 4-CF3, etc.), obtained by reaction between isonicotinoylhydrazones and diazomethane, have been prepd. and tested for antimycobacterial and anti-HIV activities. Both classes of derivs. showed interesting growth inhibitory activity on nontubercular mycobacteria, including the emerging M. avium. Such activity appears to be linked to fluorine and/or chlorine presence on the benzene rings. In contrast, none of the compds. submitted to the anti-AIDS in vitro screening displayed any protection against HIV-1 virus-induced cytopathic effect in T4-lymphocyte cell lines.

IT93-47-0

RL: RCT (Reactant)

(prepn. of halogenated isoniazid derivs. as antimycobacterial and anti-HIV agents)

RN 93-47-0 CA

4-Pyridinecarboxylic acid, [(3,4-dimethoxyphenyl)methylene]hydrazide CN (9CI)

(CA INDEX NAME)

L6 ANSWER 6 OF 7 CA COPYRIGHT 2002 ACS ACCESSION NUMBER:

TITLE:

AUTHOR(S):

117:225742 CA

Halogenated isoniazid derivatives as possible antitubercular and antineoplastic agents. Note 1 Vigorita, Maria Gabriella; Basile, Maria; Zappala, Carmela; Gabbrielli, Gabriele; Pizzimenti, Francesco

CORPORATE SOURCE:

Fac. Farm., Univ. Messina, Messina, 98168, Italy Farmaco (1992), 47(6), 893-906

SOURCE:

CODEN: FRMCE8

DOCUMENT TYPE:

LANGUAGE:

Journal English

GΙ

AΒ Isonicotinic acid arylhydrazones (I; R = Cl, F, or H; R1 = Cl, F, CF3, or OMe) were prepd. by reaction of isonicotinic acid hydrazine and the appropriate halogen-substituted benzaldehydes. Reaction of I with excess mercaptoacetic acid in refluxing anhyd. PhMe gave the corresponding 2-aryl-1,3-thiazolidin-4-ones (II), which, in turn, when oxidized by KMn04

in HOAc, gave the corresponding sulfones (III). I, II, and III were tested in vitro for antibacterial (esp. antimycobacterial), fungicidal, and antitumor (screening against 60 tumor cell lines) activities. None showed marked antimicrobial effects. Compds. bearing 3-fluoro or 3-chlorophenyl substituents had selective inhibitory effects against non-small-cell lung cancer, and those with p-Ph substituents had selective antileukemic properties. Some correlations between structure and antitumor activity are discussed.

ΙT 86189-87-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. and reaction with mercaptoacetic acid and pharmacol. of)

RN 86189-87-9 CA

CN4-Pyridinecarboxylic acid, [(4-fluorophenyl)methylene]hydrazide (9CI) (CA

INDEX NAME)

ANSWER 7 OF 7 CA COPYRIGHT 2002 ACS ACCESSION NUMBER: 89:146730 CA

TITLE:

Antimycobacterial agents. Part I.

Synthesis of some isoniazide derivatives and related

compounds

AUTHOR(S):

Umar, Muhammad; Alam, Mahbub

CORPORATE SOURCE: SOURCE:

Inst. Chem., Punjab Univ., Lahore, Pak.

Indian Chem. J. (1978), 12(12), 16-19 CODEN: ICLJAG; ISSN: 0019-4514

DOCUMENT TYPE:

Journal English

LANGUAGE:

GΙ

CH = NNHCONNHCO ОН II III

AB Hydrazone I (R = H) was prepd. by condensation of p-H2NC6H4CHO with isonicotinic acid hydrazide; I (R = Ac, EtCO) were prepd. by acylation of I (R = H). Condensation of Me salicylate and isonicotinic hydriazide gave

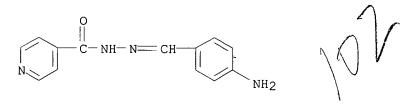
P-RNHC6H4CONHNHR1 (R1 = 4-pyridyl, R = H, Ac), III, p-RNHC6H4CH:NNHCSNH2 and 3,4,5-(MeO)3C6H2CH:NNHCOR1 were also prepd.

ΙT 6419-33-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. and acylation of)

RN 6419-33-6 CA

CN 4-Pyridinecarboxylic acid, [(4-aminophenyl)methylene]hydrazide (9CI) (CA INDEX NAME)



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FILE 'REGISTRY' ENTERED AT 03:10:29 ON 28 JAN 2002 L1STRUCTURE UPLOADED

L2 46 S L1 L3 1009 S L1 FULL

FILE 'CA' ENTERED AT 03:11:49 ON 28 JAN 2002

L4 454 S L3

L5 445 S L4 AND PD < NOVEMBER 2000

L6 7 S L5 AND ANTIMYCOBACTERIAL?

=> file caold

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FILE COVERS 1907-1966 FILE LAST UPDATED: 01 May 1997 (19970501/UP)

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L1 STRUCTURE UPLOADED

L2 46 S L1

L3 1009 S L1 FULL

FILE 'CA' ENTERED AT 03:11:49 ON 28 JAN 2002

L4 454 S L3

L5 445 S L4 AND PD < NOVEMBER 2000

L6 7 S L5 AND ANTIMYCOBACTERIAL?

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=> s 13

L7 270 L3

=> d 17, all, 1-4

- L7 ANSWER 1 OF 270 CAOLD COPYRIGHT 2002 ACS
- AN CA65:16794g CAOLD
- TI detn. of isoniazid and aconiazid by titration with N-bromosuccinimide
- AU Radecka, Czeslawa; Nigam, I. C.
- IT 54-85-3 128-08-5 **13410-86-1**
- L7 ANSWER 2 OF 270 CAOLD COPYRIGHT 2002 ACS
- AN CA65:14294b CAOLD
- TI streptomycin electroaerosols
- AU Vlasov, A. I.; Potravnova, R. S.; Naumov, G. P.; Eidel'shtein, S. I.
- IT 2779-55-7
- L7 ANSWER 3 OF 270 CAOLD COPYRIGHT 2002 ACS
- AN CA65:11321b CAOLD
- TI circular thin-layer chromatography of aromatic and .alpha.,.beta.-unsatd. aldehydes
- AU Hashmi, Manzur-ul H.; Shahid, M. A.
- IT **93-47-0** 120-57-0 **149-17-7 495-84-1** 735-97-7 4813-11-0 6956-53-2 **13059-77-3** 13059-78-4 97103-27-0
- L7 ANSWER 4 OF 270 CAOLD COPYRIGHT 2002 ACS
- AN CA65:10906e CAOLD
- TI female sex hormones in complex chemotherapy of tuberculosis
- AU Bonashevskaya, T. I.
- IT 149-17-7

=> file req

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	ENTRY	SESSION
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-4.13

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=> e 13410-86-1/rn
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E1	1	13410-82-7/RN
E2	1	13410-84-9/RN
E3	1	> 13410-86-1/RN
E4	1	13410-90-7/RN
E5	1	13410-91-8/RN
E6	1	13410-92-9/RN
E7	1	13410-93-0/RN
E8	1	13410-94-1/RN
E9	1	13410-95-2/RN
E10	1	13410-96-3/RN
E11	1	13410-98-5/RN
E12	1	13410-99-6/RN

=> s e3

L8 1 13410-86-1/RN

=> d 18

```
L8 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
```

CN 4-Pyridinecarboxylic acid, [[2-(carboxymethoxy)phenyl]methylene]hydrazide (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Isonicotinic acid, [o-(carboxymethoxy)benzylidene]hydrazide (6CI, 7CI, 8CI)

OTHER NAMES:

- CN 1-[2-(Carboxymethoxy)benzylidene]-2-isonicotinoylhydrazine
- CN Aconiazide
- CN Cpd 377
- CN Isonicophen

RN 13410-86-1 REGISTRY

```
CN Phenoxalid
FS 3D CONCORD
MF C15 H13 N3 O4
LC STN Files: BEILSTEIN*, BIOSIS, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, DDFU, DRUGNL, DRUGU, DRUGUPDATES, EMBASE, IPA, MEDLINE,
PROMT,
RTECS*, TOXCENTER, TOXLIT, USAN, USPATFULL
(*File contains numerically searchable property data)
Other Sources: WHO
```

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6 REFERENCES IN FILE CA (1967 TO DATE)
6 REFERENCES IN FILE CAPLUS (1967 TO DATE)
3 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

2779-55-7/rn Ε1 1 2779-49-9/RN E2 1 2779-54-6/RN E3 --> 2779-55-7/RN E41 2779-57-9/RN E5 1 2779-58-0/RN 2779-59-1/RN E6 1 E7 2779-60-4/RN 1 2779-62-6/RN E8 1 E9 1 2779-65-9/RN E10 1 2779-66-0/RN E11 1 2779-69-3/RN E12 2779-73-9/RN => s e3 L9 1 2779-55-7/RN => d 19

```
OTHER NAMES:
```

CN 1-(2-Carboxy-3,4-dimethoxybenzylidene)-2-isonicotinoylhydrazine

CN Benzoic acid, 2,3-dimethoxy-6-[[(4-pyridinylcarbonyl)hydrazono]methyl]-

CN Opiniazide

CN Saluside

CN Saluzid

CN Saluzide

FS 3D CONCORD

MF C16 H15 N3 O5

CI COM

LC STN Files: ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, CA, CAOLD, CAPLUS, CHEMCATS, DDFU, DRUGU, EMBASE, IPA, MEDLINE, MRCK*, RTECS*, TOXCENTER, TOXLIT, USAN

(*File contains numerically searchable property data) Other Sources: $$\operatorname{WHO}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 63 REFERENCES IN FILE CA (1967 TO DATE)
- 6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 63 REFERENCES IN FILE CAPLUS (1967 TO DATE)
- 27 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

	^	149-17-7/	~~
=>	e	149-1/-//	rn

E1	1		149-15-5/RN
E2	1		149-16-6/RN
E3	1	>	149-17-7/RN
E4	1		149-19-9/RN
E5	1		149-20-2/RN
E6	1		149-21-3/RN
E7	1		149-22-4/RN
E8	1		149-23-5/RN
E9	1		149-24-6/RN
E10	1		149-26-8/RN
E11	1		149-29-1/RN
E12	1		149-30-4/RN

=> s e3

L10 1 149-17-7/RN

=> d 110

```
L10 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
     149-17-7 REGISTRY
RN
     4-Pyridinecarboxylic acid,
CN
[(4-hydroxy-3-methoxyphenyl)methylene]hydrazide
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Isonicotinic acid, vanillylidenehydrazide (6CI, 8CI)
OTHER NAMES:
CN
     Ftivazid
CN
     Ftivazide
CN
     N-Isonicotinamido-3-methoxy-4-hydroxybenzalimine
CN
     Phthivazid
CN
     Phthivazide
CN
     Vanicid
CN
     Vanillaberon
CN
     Vanizide
     3D CONCORD
FS
MF
     C14 H13 N3 O3
CI
     COM
LC
                  ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, CA, CANCERLIT,
     STN Files:
       CAOLD, CAPLUS, CASREACT, CHEMCATS, DDFU, DRUGU, EMBASE, GMELIN*, IPA,
       MEDLINE, RTECS*, SPECINFO, TOXCENTER, TOXLIT, USAN
         (*File contains numerically searchable property data)
     Other Sources:
                      WHO
```

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

128 REFERENCES IN FILE CA (1967 TO DATE)
11 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
128 REFERENCES IN FILE CAPLUS (1967 TO DATE)
100 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> d his

L1

(FILE 'HOME' ENTERED AT 03:10:17 ON 28 JAN 2002)

FILE 'REGISTRY' ENTERED AT 03:10:29 ON 28 JAN 2002 STRUCTURE UPLOADED

L2 . 46 S L1

L3 1009 S L1 FULL

FILE 'CA' ENTERED AT 03:11:49 ON 28 JAN 2002 . L4 454 S L3 L5445 S L4 AND PD < NOVEMBER 2000 L6 7 S L5 AND ANTIMYCOBACTERIAL? FILE 'CAOLD' ENTERED AT 03:14:20 ON 28 JAN 2002 L7 270 S L3 FILE 'REGISTRY' ENTERED AT 03:14:50 ON 28 JAN 2002 E 13410-86-1/RN L8 1 S E3 E 2779-55-7/RN L9 1 S E3 E 149-17-7/RN L10 1 S E3 => ---Logging off of STN---

=>
Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	6.34	183.27
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY 0.00	SESSION -4.13

STN INTERNATIONAL LOGOFF AT 03:17:32 ON 28 JAN 2002

Connection closed by remote host